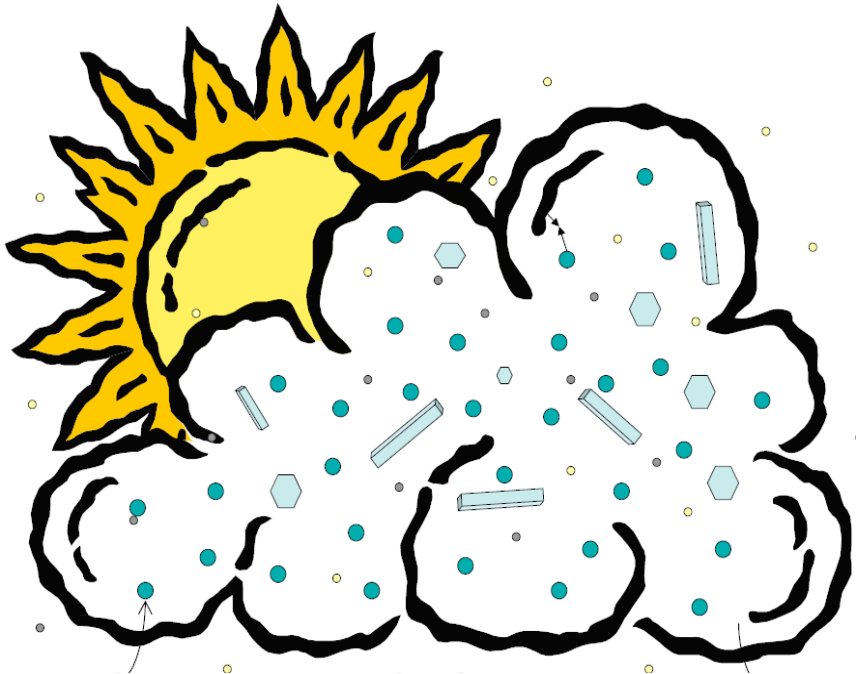


Indirect and Semi-Direct Aerosol Campaign

PI: Steve Ghan, Pacific Northwest National Laboratory

Co-I: Sarah Brooks, Don Collins, Connor Flynn, Ann Fridlind, Tim Garrett, John Hallett, John Hubbe, Greg Kok, Dan Lubin, Greg McFarquhar, David Mitchell, Mike Poellet, Matthew Shupe, David Turner, Hans Verlinde, Shaocheng Xie



Barrow, Alaska

April 2008



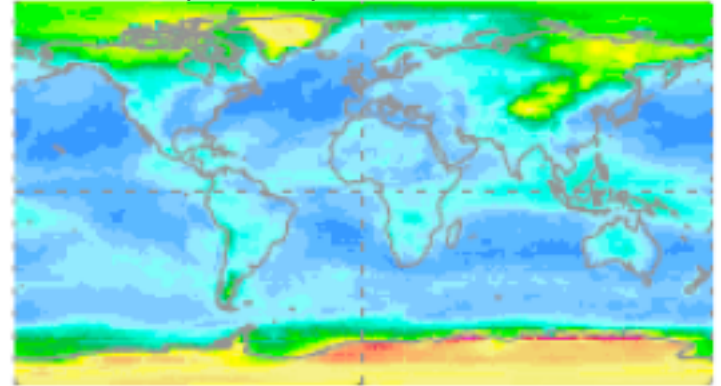
ISDAC Motivation

- Most studies of cloud-aerosol interactions have focused on warm clouds.
- Cloud-aerosol interactions are much more complex for ice or mixed-phase clouds than for warm clouds.
- The Mixed-Phase Arctic Cloud Experiment at the ARM site at Barrow in October 2004 has provided new insight into these interactions.
- The arctic air during April is expected to be much more polluted than the air during M-PACE.
- This contrast provides an opportunity to
 - distinguish between aerosol effects on arctic clouds under clean and polluted conditions
 - improve understanding of the scavenging of arctic aerosol during spring
 - identify the chemical signature of ice nuclei in the arctic

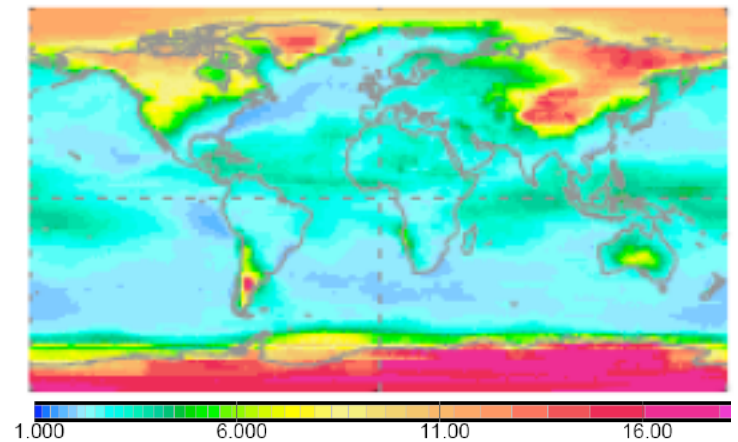
Why Study Scavenging in the Arctic?

- Most of the relative uncertainty in simulated aerosol optical depth and mass loading is in polar regions.
- Most arctic aerosol comes from midlatitude sources.
- The treatment of transport is unlikely to cause a 10-fold uncertainty.
- Such uncertainty is probably due to the treatment of scavenging by clouds.

Max/Min of Central 2/3 of 16 Models
Aerosol Optical Depth



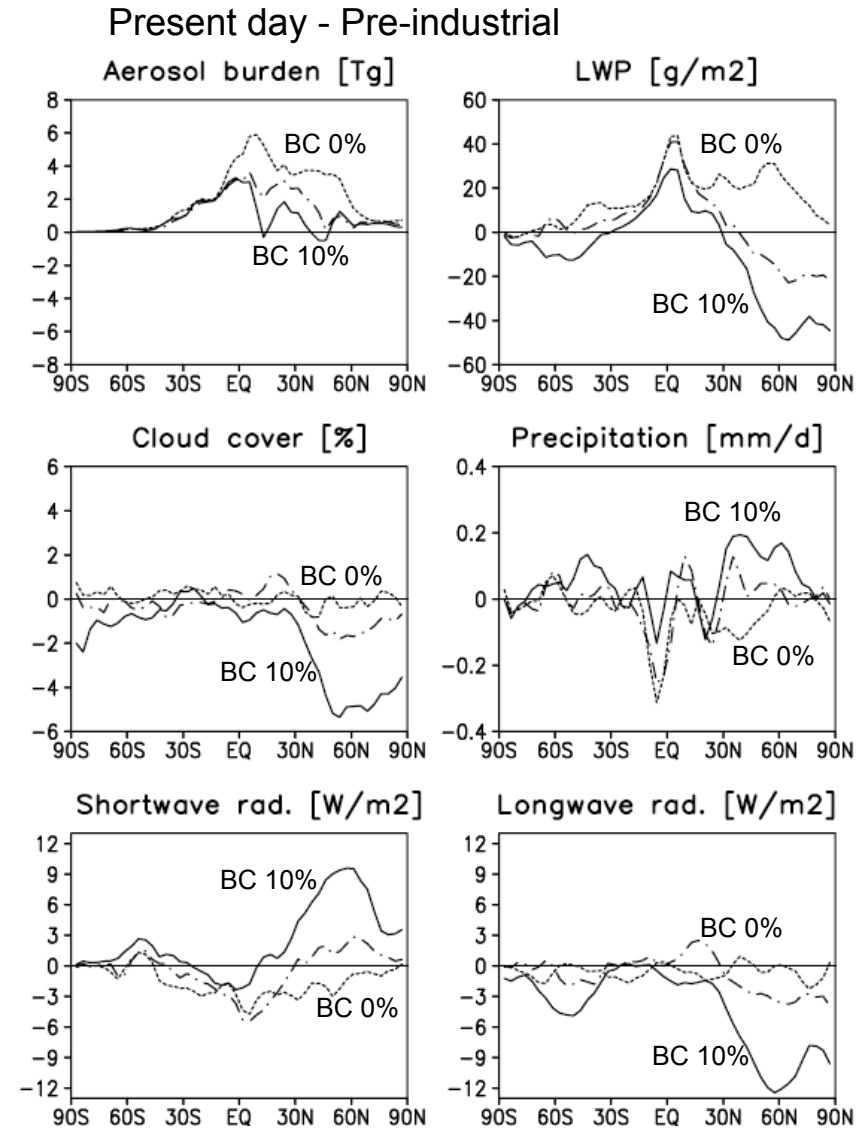
Aerosol Column Mass



Kinne et al., An AeroCom initial assessment.
Atmos. Chem. & Phys., 2006.

Why Study Ice Nucleation in the Arctic?

- Anthropogenic changes in ice nucleation may influence aerosol burden and climate.
- The treatment of ice nucleation in models affects the aerosol scavenging efficiency.
- A variety of ice nucleation mechanisms are involved.
- A counterflow virtual impactor can be used to isolate ice nuclei from the cloud particles and then determine size and composition.
- ASP could provide the first Arctic measurements of ice nuclei size and composition.



Lohmann, A glaciation indirect aerosol effect caused by soot aerosols. GRL 2002.

Key ARM Aircraft Measurements

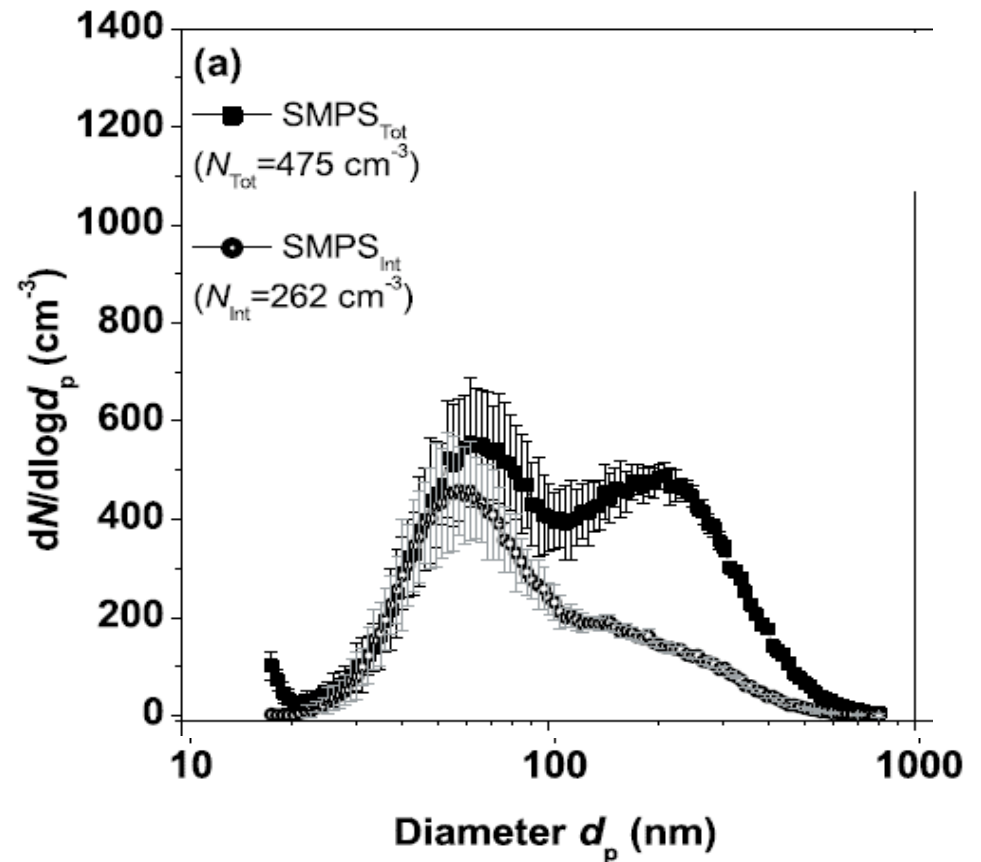
Instrument	Measurements
TSI 3025	total particle concentration ($> 3 \text{ nm}$)
DMA	aerosol size distribution ($0.01\text{-}0.75 \text{ }\mu\text{m}$)
PCASP	aerosol size distribution ($0.1\text{-}3 \text{ }\mu\text{m}$)
TDMA	Size-resolved aerosol hygroscopicity ($0.015 - 0.6 \text{ }\mu\text{m}$)
DMT CCN counter	CCN concentration (one S)
CFDC	IN concentration
PSAP	optical absorption
Gust probe	updraft velocity
DMT CAPS	temperature, LWC, cloud particle size dist ($0.5\text{-}1500 \text{ }\mu\text{m}$)
DMT CSI	total condensed water concentration
SPEC CPI	cloud particle image $15\text{-}2500 \text{ }\mu\text{m}$
CIN	cloud extinction coefficient, asymmetry parameter

ASP Instruments and Measurements

Instrument	Measurement
Counterflow Virtual Impactor	Cloud-borne aerosol
Scanning Mobility Particle Sizer	Aerosol size distribution 3-1000 nm
PCASP	Aerosol size distribution 0.1-3 μm
TSI 3010, 3025A	Total aerosol number
DRI CCN Spectrometer	CCN spectrum
Particle-in-Liquid System	Particle ionic composition
Aerosol Mass Spectrometer	Size-resolved composition
Time-Resolved Aerosol Collector / CCSEM/EDX	Single particle chemical composition and mixing state
DRI Photoacoustic	Aerosol absorption

Aerosol Scavenging

- Two conditions for wet scavenging of aerosol:
 - Attachment to hydrometeor
 - Precipitation of hydrometeor
- Evaluate first condition by comparing simulated and observed partitioning of aerosol between interstitial and cloud-borne
- Evaluate second by comparing simulated and observed hydrometeor size distribution and precipitation rate



Henning, Bojinski, Diehl, Ghan, Nyeki, Weingartner, Wurzler, and Baltensperger: Aerosol partitioning in natural mixed-phase clouds. GRL 2004.

Deep Convective Clouds and Chemistry

June-July 2009 Colorado - Oklahoma



Steven Ghan (PNNL)

Mary Barth, Bill Brune, Chris Cantrell (NCAR)

Steve Rutledge (CSU)

- Deep convection plays important roles in vertical transport, aqueous-phase sulfate production, nucleation, and scavenging of trace chemicals and particulate matter.
- The representation of this influence in global climate models is highly uncertain.
- Several ASP projects address this challenge.
- DOE does not have an aircraft that can sample detrainment of trace species from deep convection.
- DC3 would provide upper tropospheric measurements that would complement DOE measurements in the lower troposphere.

Pollutant Mass Closure (Don Lenschow)

- Circles around cloud at 3 km altitude intervals

- Flux of q into cloud: $F_q(z) = \frac{1}{A} \oint_C v_{\perp} q dl$

- Entrainment below z_1 : $E_q = \int_0^{z_1} \rho F_q dz$

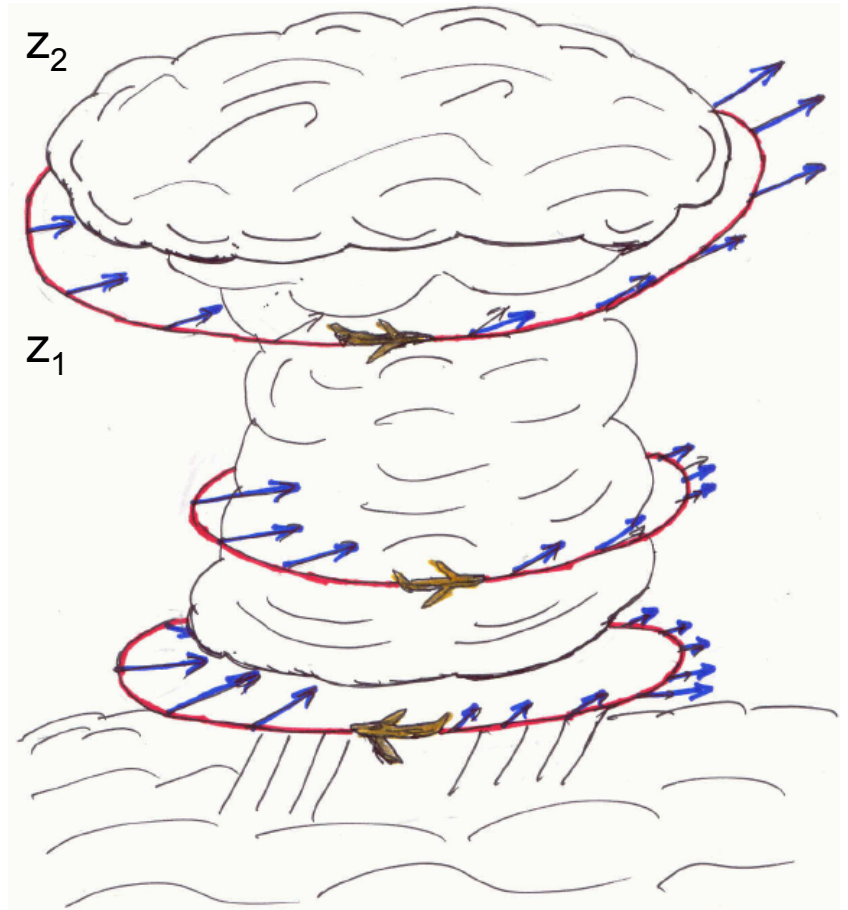
- Detrainment above z_1 : $D_q(z) = - \int_{z_1}^z \rho F_q dz'$

- Scavenging ratio $S_q = \frac{E_q - D_q}{E_q}$

- $S=0$ for dry air

- $S \sim 1$ for water, soluble gases and large soluble particles

- $S \ll 1$ for insoluble gases



Key Platforms

- DC3 would provide
 - NSF/NCAR G-V to sample the convective *outflow* of aerosol and trace gases in the upper troposphere
 - either the NASA DC-8 or the NSF/NCAR C-130 to sample the *inflow* of aerosol and trace gases in the middle troposphere
- ASP would provide the G-1 to sample the *inflow* of aerosol and trace gases in the lower troposphere

Key Measurements

Measurement	DC-8	G-V	G-1
O ₃ , CO, HNO ₃ , SO ₂	1	1	1
NO, NO _y , NO ₂	1	1	1
H ₂ O vapor	1	1	1
Peroxides	1	1	1
CVI		2	
Aerosol size distribution	1	2	1
CCN	1	2	1
Size-resolved aerosol composition (AMS, PILS)	1	2	1
Time-Resolved Aerosol Collector		2	1
Winds, T, p, location	1	1	1

1: essential. 2: desirable